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=> s antidepressant? and ((rapid or immediate)(w)release)
L1 692 ANTIDEPRESSANT? AND ((RAPID OR IMMEDIATE)(W) RELEASE)

=> s l1 and (sexual dysfunction)

L2 59 L1 AND (SEXUAL DYSFUNCTION)

=> s 12 and (premature ejaculation)

L3 12 L2 AND (PREMATURE EJACULATION)

=> d 13 1-12 ibib abs

L3 ANSWER 1 OF 12 USPATFULL on STN

ACCESSION NUMBER:

2004:139435 USPATFULL

TITLE:

N-phenpropylcyclopentyl-substituted glutaramide derivatives as inhibitors of neutral endopeptidase

INVENTOR(S):

Challenger, Stephen, Sandwich, UNITED KINGDOM
Cook, Andrew Simon, Sandwich, UNITED KINGDOM
Gillmore, Adam Thomas, Sandwich, UNITED KINGDOM
Middleton, Donald Stuart, Sandwich, UNITED KINGDOM
Pryde, David Cameron, Sandwich, UNITED KINGDOM

Stobie, Alan, Sandwich, UNITED KINGDOM

PATENT ASSIGNEE(S): Pfizer Inc (non-U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 2004106611 A1 20040603
APPLICATION INFO.: US 2003-696021 A1 20031028 (10)

RELATED APPLN. INFO.: Division of Ser. No. US 2002-96218, filed on 12 Mar

2002, GRANTED, Pat. No. US 6660756

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: PFIZER INC., PATENT DEPARTMENT, MS8260-1611, EASTERN

POINT ROAD, GROTON, CT, 06340

NUMBER OF CLAIMS:

EXEMPLARY CLAIM: 1
NUMBER OF DRAWINGS: 1

1 Drawing Page(s)

LINE COUNT:

4453

28

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The invention relates to compounds of formula (I) for treating for example sexual dysfunction, wherein R.sup.1 is optionally substituted C.sub.1-6alkyl, optionally substituted carbocyclyl, optionally substituted heterocyclyl, hydrogen, C.sub.1-6alkoxy, --NR.sup.2R.sup.3 or --NR.sup.4SO.sub.2R.sup.5; X is the linkage -- (CH.sub.2).sub.n-- or -- (CH.sub.2).sub.q--O-- (wherein Y is attached to the oxygen); wherein one or more hydrogen atoms in linkage X may be replaced independently by C.sub.1-4alkoxy; hydroxy; hydroxy(C.sub.1-3alkyl); C.sub.3-7cycloalkyl; carbocyclyl; heterocyclyl; or by C.sub.1-4alkyl optionally substituted by one or more fluoro or phenyl groups; n is 3, 4, 5, 6 or 7; and q is 2, 3, 4, 5 or 6; and Y is phenyl or pyridyl, each of which may be substituted; or two R.sup.8 groups on adjacent carbon atoms together with the interconnecting carbon atoms may form a fused optionally substituted 5- or 6-membered carbocyclic or heterocyclyic ring. ##STR1##

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 2 OF 12 USPATFULL on STN

ACCESSION NUMBER:

2004:101836 USPATFULL

TITLE:

Ligands for monoamine receptors and transporters, and

methods of use thereof

INVENTOR(S):

Aquila, Brian M., Marlborough, MA, UNITED STATES Bannister, Thomas D., Northborough, MA, UNITED STATES Cuny, Gregory D., Somerville, MA, UNITED STATES Hauske, James R., Concord, MA, UNITED STATES Holland, Joanne M., Brookline, MA, UNITED STATES

Persons, Paul E., Westborough, MA, UNITED STATES Radeke, Heike, S. Grafton, MA, UNITED STATES Wang, Fengjiang, Northborough, MA, UNITED STATES

Shao, Liming, Lincoln, MA, UNITED STATES

KIND DATE

PATENT INFORMATION: US 2004077706 A1 20040422 APPLICATION INFO.: US 2003-607457 A1 20030626 (10)

RELATED APPLN. INFO.: Division of Ser. No. US 2001-951130, filed on 12 Sep

2001, PENDING

NUMBER DATE

PRIORITY INFORMATION:

US 2001-273530P 20010305 (60) US 2001-298057P 20010613 (60)

Utility DOCUMENT TYPE:

APPLICATION

LEGAL REPRESENTATIVE: FOLEY HOAG, LLP, PATENT GROUP, WORLD TRADE CENTER WEST,

FOLEY HOAG, LLP, PATENT GROUP, WORLI
155 SEAPORT BLVD, BOSTON, MA, 02110

NUMBER OF CLAIMS: 172

EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 1 DrawiLINE COUNT:
CAS TOTAL

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

One aspect of the present invention relates to heterocyclic compounds. A second aspect of the present invention relates to the use of the heterocyclic compounds as ligands for various mammalian cellular receptors, including dopamine, serotonin, or norepinephrine transporters. The compounds of the present invention will find use in the treatment of numerous ailments, conditions and diseases which afflict mammals, including but not limited to addiction, anxiety, depression, sexual dysfunction, hypertension, migraine, Alzheimer's disease, obesity, emesis, psychosis, schizophrenia, Parkinson's disease, inflammatory pain, neuropathic pain, Lesche-Nyhane disease, Wilson's disease, and Tourette's syndrome. An

additional aspect of the present invention relates to the synthesis of combinatorial libraries of the heterocyclic compounds, and the screening of those libraries for biological activity, e.g., in assays based on dopamine transporters.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 3 OF 12 USPATFULL on STN

ACCESSION NUMBER:

2003:153464 USPATFULL

TITLE:

N-phenpropylcuclopentyl-substituted glutaramide derivatives as inhibitors of neutral endopeptidase

INVENTOR(S):

Challenger, Stephen, Sandwich, UNITED KINGDOM Cook, Andrew Simon, Sandwich, UNITED KINGDOM Gillmore, Adam Thomas, Sandwich, UNITED KINGDOM Middleton, Donald Stuart, Sandwich, UNITED KINGDOM

)

Pryde, David Cameron, Sandwich, UNITED KINGDOM

Stobie, Alan, Sandwich, UNITED KINGDOM

	NUMBER	KIND	DATE	
PATENT INFORMATION:	US 2003105132	A1	20030605	
APPLICATION INFO.:	US 6660756 US 2002-96218	B2 A1	20031209 20020312	(10)
	00 000 00010		20020312	(- 0)

		NUMBER	DATE	
PRIORITY	INFORMATION:	GB 2001-7750	20010328	
	(GB 2001-13112	20010530	
		GB 2001-20152	20010817	
		US 2001-292485P	20010521	(60)
		US 2001-299031P	20010618	(60)
		US 2001-317777P	20010906	(60)

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: Gregg C. Benson, Pfizer Inc., Patent Department, MS

4159, Eastern Point Road, Groton, CT, 06340

NUMBER OF CLAIMS: 28 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 1 Drawing Page(s) LINE COUNT: 4350

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The invention relates to compounds of formula (I) for treating for example sexual dysfunction, wherein R.sup.1 is optionally substituted C.sub.1-6alkyl, optionally substituted carbocyclyl, optionally substituted heterocyclyl, hydrogen, C.sub.1-6alkoxy, --NR.sup.2 R.sup.3 or --NR.sup.4SO.sub.2R.sup.5; X is the linkage -- (CH.sub.2).sub.n-- or -- (CH.sub.2).sub.q--O-- (wherein Y is attached to the oxygen); wherein one or more hydrogen atoms in linkage X may be replaced independently by C.sub.1-4alkoxy; hydroxy; hydroxy(C.sub.1-3alkyl); C.sub.3-7cycloalkyl; carbocyclyl; heterocyclyl; or by C.sub.1-4alkyl optionally substituted by one or more fluoro or phenyl groups; n is 3, 4, 5, 6 or 7; and q is 2, 3, 4, 5 or 6; and Y is phenyl or pyridyl, each of which may be substituted; or two R.sup.8 groups on adjacent carbon atoms together with the interconnecting carbon atoms may form a fused optionally substituted 5- or 6-membered carbocyclic or heterocyclyic ring. ##STR1##

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 4 OF 12 USPATFULL on STN

ACCESSION NUMBER: 2003:153429 USPATFULL Alkylamide compounds

INVENTOR(S): Simon, Andrew, Sandwich, UNITED KINGDOM

Stobie, Alan, Sandwich, UNITED KINGDOM

PATENT ASSIGNEE(S): Pfizer Inc. (non-U.S. corporation)

NUMBER KIND DATE _____ PATENT INFORMATION: US 2003105097 A1 20030605 APPLICATION INFO.: US 2002-139443 A1 20020506 (10)

NUMBER DATE

PRIORITY INFORMATION: GB 2001-11709 20010514 US 2001-294278P 20010530 (60)

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: PFIZER INC, 150 EAST 42ND STREET, 5TH FLOOR - STOP 49,

NEW YORK, NY, 10017-5612

NUMBER OF CLAIMS: 11
EXEMPLARY CLAIM: 1
2001

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The invention provides compounds of formula (I) ##STR1##

wherein R.sup.1 is optionally substituted C.sub.1-6alkyl, optionally substituted carbocyclyl, optionally substituted heterocyclyl, C.sub.1-6alkoxy, --NR.sup.2R.sup.3 or --NR.sup.4SO.sub.2R.sup.5; n is 2 to 6; X is oxygen, sulfur or --CH.sub.2--; Y is C.sub.1-6alkyl which may be branched- or straight-chain, and may be independently substituted by one or more halo, C.sub.1-4alkoxy, hydroxy or C.sub.3-7cycloalkyl; wherein the linkage -- (CH.sub.2).sub.n-- and the linkage when X is --CH.sub.2--, may be substituted by C.sub.1-4alkyl, C.sub.1-4haloalkyl, hydroxy, C.sub.1-4alkoxy, C.sub.1-4haloalkoxy, hydroxyC.sub.1-4alkyl, C.sub.1-4alkoxyC.sub.1-4alkyl, C.sub.1-4haloalkoxyC.sub.1-4alkyl, C.sub.3-7cycloalkyl or C.sub.3-7cycloalkylC.sub.1-4alkyl. The compounds of the invention are useful in the treatment of sexual dysfunction, particularly female sexual dysfunction.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 5 OF 12 USPATFULL on STN

ACCESSION NUMBER: 2003:153403 USPATFULL

TITLE: Thiazole and other heterocyclic ligands for mammalian

dopamine, muscarinic and serotonin receptors and

transporters, and methods of use thereof

INVENTOR(S): Cuny, Gregory D., Somerville, MA, UNITED STATES

Hauske, James R., Concord, MA, UNITED STATES

Heffernan, Michele L.R., Worcester, MA, UNITED STATES

Holland, Joanne M., Brookline, MA, UNITED STATES

Persons, Paul E., Westborough, MA, UNITED STATES

Radeke, Heike, Grafton, MA, UNITED STATES

NUMBER KIND DATE ______ US 2003105071 A1 20030605 US 6699866 B2 20040302 US 2002-123089 A1 20020412 (10) PATENT INFORMATION: APPLICATION INFO.:

NUMBER DATE _____

PRIORITY INFORMATION: US 2001-284159P 20010417 (60) US 2001-313648P 20010820 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FOLEY HOAG, LLP, PATENT GROUP, WORLD TRADE CENTER WEST,

155 SEAPORT BLVD, BOSTON, MA, 02110

106 NUMBER OF CLAIMS: EXEMPLARY CLAIM: LINE COUNT: 5332

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

One aspect of the present invention relates to novel heterocyclic compounds. A second aspect of the present invention relates to the use of the novel heterocyclic compounds as ligands for various mammalian cellular receptors, including G-protein coupled receptors. A third aspect of the present invention relates to the use of the novel heterocyclic compounds as ligands for mammalian dopamine, muscarinic or serotonin receptors or transporters. Another aspect of the present invention relates to the use of the novel heterocyclic compounds as ligands for mammalian dopamine, muscarinic or serotonin receptors. The compounds of the present invention will also find use in the treatment of numerous ailments, conditions and diseases which afflict mammals, including but not limited to addiction, anxiety, depression,

sexual dysfunction, hypertension, migraine,

Alzheimer's disease, obesity, emesis, psychosis, analgesia, schizophrenia, Parkinson's disease, restless leg syndrome, sleeping disorders, attention deficit hyperactivity disorder, irritable bowel syndrome, premature ejaculation, menstrual dysphoria syndrome, urinary incontinence, inflammatory pain, neuropathic pain, Lesche-Nyhane disease, Wilson's disease, Tourette's syndrome, psychiatric disorders, stroke, senile dementia, peptic ulcers, pulmonary obstruction disorders, and asthma.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 6 OF 12 USPATFULL on STN

ACCESSION NUMBER: 2003:72022 USPATFULL

TITLE: Ligands for monoamine receptors and transporters, and

methods of use thereof

INVENTOR(S): Aquila, Brian M., Marlborough, MA, UNITED STATES

Bannister, Thomas D., Northborough, MA, UNITED STATES

Cuny, Gregory D., Somervill, MA, UNITED STATES Hauske, James R., Concord, MA, UNITED STATES Holland, Joanne M., Brookline, MA, UNITED STATES Persons, Paul E., Westborough, MA, UNITED STATES Radeke, Heike, S. Grafton, MA, UNITED STATES Wang, Fengjiang, Northborough, MA, UNITED STATES

Shao, Liming, Lincoln, MA, UNITED STATES

NUMBER	KIND	DATE	
US 2003050309 US 2001-951130	A1 A1	20030313	(9)

			NUMBER	DATE	
PRIORITY	INFORMATION:	US	2000-231667P	20000911	(60)
		US	2001-273530P	20010305	(60)
		US	2001-298057P	20010613	(60)
DOCUMENT	MYDH.	TT4 -			

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FOLEY HOAG LLP, PATENT GROUP, 155 SEAPORT BOULEVARD,

BOSTON, MA, 02110

NUMBER OF CLAIMS: 172 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 1 Drawing Page(s)

LINE COUNT: 8267

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

One aspect of the present invention relates to heterocyclic compounds. A second aspect of the present invention relates to the use of the heterocyclic compounds as ligands for various mammalian cellular

receptors, including dopamine, serotonin, or norepinephrine transporters. The compounds of the present invention will find use in the treatment of numerous ailments, conditions and diseases which afflict mammals, including but not limited to addiction, anxiety, depression, sexual dysfunction, hypertension, migraine, Alzheimer's disease, obesity, emesis, psychosis, schizophrenia, Parkinson's disease, inflammatory pain, neuropathic pain, Lesche-Nyhane disease, Wilson's disease, and Tourette's syndrome. An additional aspect of the present invention relates to the synthesis of combinatorial libraries of the heterocyclic compounds, and the screening of those libraries for biological activity, e.g., in assays based on dopamine transporters.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 7 OF 12 USPATFULL on STN

ACCESSION NUMBER: 2002:332475 USPATFULL

TITLE: On demand administration of clomipramine and salts

thereof to treat premature

ejaculation

INVENTOR(S): Tam, Peter, Redwood City, CA, United States

> Gesundheit, Neil, Los Altos, CA, United States Wilson, Leland F., Menlo Park, CA, United States

PATENT ASSIGNEE(S): Vivus Inc., Mountain View, CA, United States (U.S.

corporation)

NUMBER KIND DATE -----

PATENT INFORMATION: US 6495154 B1 20021217 APPLICATION INFO.: US 2000-721412 20001121 (9) DOCUMENT TYPE: Utility

FILE SEGMENT: GRANTED
PRIMARY EXAMINER: Azpuru, Carlos

LEGAL REPRESENTATIVE: Reed & Associates, Reed, Dianne E.

NUMBER OF CLAIMS: 71 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 0 Drawing Figure(s); 0 Drawing Page(s)

LINE COUNT: 1221

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

A method is provided for delaying the onset of ejaculation in an individual. The method involves systemic and on demand administration to an individual of a pharmaceutical formulation containing an amount of an active agent selected from the group consisting of clomipramine and pharmacologically acceptable acid addition salts thereof. Drug delivery may be accomplished via any route designed to provide systemic levels of the active agent effective to delay the onset of ejaculation. Pharmaceutical formulations and dosage forms are provided as well.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 8 OF 12 USPATFULL on STN

ACCESSION NUMBER: 2002:288146 USPATFULL

TITLE: As-needed administration of tricyclic and other non-SRI

antidepressant drugs to treat premature

ejaculation

INVENTOR(S): Tam, Peter, Redwood City, CA, UNITED STATES

> Gesundheit, Neil, Los Altos, CA, UNITED STATES Wilson, Leland F., Menlo Park, CA, UNITED STATES

NUMBER KIND DATE ______ PATENT INFORMATION: US 2002161016 A1 20021031
APPLICATION INFO.: US 2001-996407 A1 20011121 (9)
RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 2000-721412, filed

on 21 Nov 2000, PENDING

DOCUMENT TYPE: Utility APPLICATION APPLICATION

LEGAL REPRESENTATIVE: REED & ASSOCIATES, 800 MENLO AVENUE, SUITE 210, MENLO

PARK, CA, 94025

NUMBER OF CLAIMS:

72

EXEMPLARY CLAIM: 1
LINE COUNT: 14

1452

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

A method is provided for treatment of premature

ejaculation by administration of an antidepressant

drug selected from tricyclic antidepressants, tetracyclic

antidepressants, MAO inhibitors, azaspirone

antidepressants, and atypical non-SRI antidepressants.

In a preferred embodiment, administration is on as "as-needed" basis, i.e., the drug is administered immediately or at most several hours prior to sexual activity. Pharmaceutical formulations and packaged kits are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 9 OF 12 USPATFULL on STN

ACCESSION NUMBER: 2002:172371 USPATFULL

TITLE:

Treatment of premature ejaculation

INVENTOR(S): Boolell, Mitradev, County of Kent, UNITED KINGDOM

NUMBER KIND DATE -----PATENT INFORMATION: US 2002091129 A1 20020711 APPLICATION INFO.: US 2001-990955 A1 20011116 (9)

NUMBER DATE ______

PRIORITY INFORMATION: GB 2000-28245 20001120 US 2001-260564P 20010109 (60)

DOCUMENT TYPE:

Utility APPLICATION

FILE SEGMENT:

LEGAL REPRESENTATIVE: Gregg C. Benson, Pfizer Inc., Patent Department, MS

4159, Eastern Point Road, Groton, CT, 06340

NUMBER OF CLAIMS: 11
EXEMPLARY CLAIM: 1
LINE COUNT: 863 LINE COUNT: 862

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

This invention relates to the use of cyclic guanosine 3',

5'-monophosphate phosphodiesterase type five inhibitors, including in particular the compound sildenafil, for the treatment of

premature ejaculation in patients with normal erectile

function.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 10 OF 12 USPATFULL on STN

ACCESSION NUMBER:

2002:67175 USPATFULL

TITLE:

Administration of phosphodiesterase inhibitors for the

treatment of premature ejaculation

INVENTOR(S):

Wilson, Leland F., Menlo Park, CA, UNITED STATES Doherty, Paul C., JR., Cupertino, CA, UNITED STATES

Place, Virgil A., Kawaihae, HI, UNITED STATES Smith, William L., Montclair, NJ, UNITED STATES Abdel-Hamid Abdou Ali, Ibrahim AbouBakr, Mansoura,

EGYPT

NUMBER KIND DATE ------

US 2002037828 A1 20020328 US 6403597 B2 20020611 US 2001-888250 A1 20010621 (9) PATENT INFORMATION:

APPLICATION INFO.:

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1999-467094, filed on 10 Dec 1999, PENDING Continuation-in-part of Ser. No. US 1998-181070, filed on 27 Oct 1998, GRANTED, Pat.

No. US 6037346 Continuation-in-part of Ser. No. US

1997-958816, filed on 28 Oct 1997, ABANDONED

Utility DOCUMENT TYPE: APPLICATION FILE SEGMENT:

LEGAL REPRESENTATIVE: REED & ASSOCIATES, 800 MENLO AVENUE, SUITE 210, MENLO

PARK, CA, 94025 94 1 NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 1 Drawing Page(s)
LINE COUNT: 2011

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

A method is provided for treatment of premature

ejaculation by administration of a phosphodiesterase inhibitor, e.g., an inhibitor of a Type III, Type IV, or Type V phosphodiesterase. In a preferred embodiment, administration is on as "as needed" basis, i.e., the drug is administered immediately or several hours prior to sexual activity. Pharmaceutical formulations and packaged kits are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 11 OF 12 USPATFULL on STN

ACCESSION NUMBER: 1998:134657 USPATFULL Low dose fluoxetine tablet TITLE:

El-Rashidy, Ragab, Deerfield, IL, United States INVENTOR(S):

Ronsen, Bruce, River Forest, IL, United States

Pentech Pharmaceuticals, Inc., Wheeling, IL, United PATENT ASSIGNEE(S):

States (U.S. corporation)

NUMBER KIND DATE -----PATENT INFORMATION: US 5830500 19981103
APPLICATION INFO.: US 1996-681276 19960722 (8)
DOCUMENT TYPE: Utility
FILE SEGMENT: Granted
PRIMARY EXAMINER: Kulkosky, Peter F.

LEGAL REPRESENTATIVE: Olson & Hierl, Ltd.

NUMBER OF CLAIMS: 8 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 1 Drawing Figure(s); 1 Drawing Page(s)

339 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

A direct compression tablet exhibiting relatively low hardness provides

a relatively rapid release of fluoxetine. The tablet

has a hardness of no more than about 6 kilopascals and a dicalcium phosphate dihydrate-to-disintegrant weight ratio of about 3 to about 7.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 12 OF 12 EUROPATFULL COPYRIGHT 2004 WILA on STN

PATENT APPLICATION - PATENTANMELDUNG - DEMANDE DE BREVET

ACCESSION NUMBER: 1258474 EUROPATFULL EW 200247 FS OS

Alkylamide compounds. TITLE: Alkylamid-Derivate. Derives d'alkylamide.

Cook, Andrew Simon, Pfizer Global Res. and Dev., INVENTOR(S):

Ramsgate Road, Sandwich, Kent CT13 9NJ, GB;

Stobie, Alan, Pfizer Global Res. and Dev., Ramsgate

Road, Sandwich, Kent CT13 9NJ, GB

Pfizer Limited, Ramsgate Road, Sandwich, Kent CT13 9NJ, PATENT ASSIGNEE(S):

GB, in GB;

PFIZER INC., 235 East 42nd Street, New York, N.Y. 10017, US, in BE, CH, DE, DK, ES, FI, FR, GR, IE, IT, LI, LU,

MC, NL, PT, SE, AT, CY

PATENT ASSIGNEE NO:

204310; 200961

AGENT:

Rutt, Jason Edward et al., Pfizer Limited European Patents Department Ramsgate Road, Sandwich, Kent CT13

9NJ, GB

AGENT NUMBER:

99591

OTHER SOURCE:

BEPA2002097 EP 1258474 A2 0036

SOURCE:

Wila-EPZ-2002-H47-T1a

DOCUMENT TYPE:

Patent

LANGUAGE: DESIGNATED STATES: Anmeldung in Englisch; Veroeffentlichung in Englisch R AT; R BE; R CH; R CY; R DE; R DK; R ES; R FI; R FR; R GB; R GR; R IE; R IT; R LI; R LU; R MC; R NL; R PT; R

SE; R TR; R AL; R LT; R LV; R MK; R RO; R SI

PATENT INFO.PUB.TYPE: EPA2 EUROPAEISCHE PATENTANMELDUNG PATENT INFORMATION:

	PATENT NO		KIND	DATE
	ΕP	1258474	A2	20021120
'OFFENLEGUNGS' DATE:				20021120
APPLICATION INFO.:	ΕP	2002-253114		20020502
PRIORITY APPLN. INFO.:	GB	2001-111709		20010514